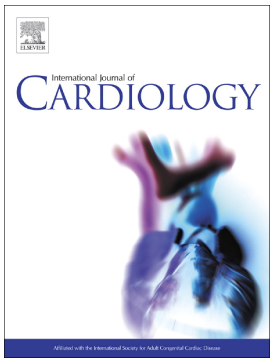
Since January 2020 Elsevier has created a COVID-19 resource centre with free information in English and Mandarin on the novel coronavirus COVID-19. The COVID-19 resource centre is hosted on Elsevier Connect, the company's public news and information website.



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PII: S0167-5273(21)01499-6

DOI: https://doi.org/10.1016/j.ijcard.2021.09.058

Reference: IJCA 29821

To appear in: *International Journal of Cardiology*

Received date: 4 July 2021

Revised date: 26 September 2021

Accepted date: 28 September 2021

Please cite this article as: A. Sisinni, L. Rossi, A. Battista, et al., Pre-admission acetylsalicylic acid therapy and impact on in-hospital outcome in COVID-19 patients: The ASA-CARE study, *International Journal of Cardiology* (2021), https://doi.org/10.1016/j.ijcard.2021.09.058

Pre-admission Acetylsalicylic Acid Therapy and Impact on In-hospital Outcome in COVID-19 patients: the ASA-CARE study.

Antonio Sisinni,[1] MD, Luca Rossi,[2] MD, Antonio Battista,[3] MD, Enrico Poletti,[4] MD, Federica Battista,[5] MD, Rosa Alessia Battista,[6] MD, Alessandro Malagoli,[7] MD, Andrea Biagi,[8] MD, Alessia Zanni,[9] MD, Concetta Sticozzi,[10] MD, Greta Comastri,[11] MD, Massimiliano M Marrocco-Trischitta,[12] MD, Alberto Monello,[13] MD, Alberto Margonato,[14] MD, Francesco Bandera,[15] MD, Pasquale Vergara,[16] MD, Marco Guazzi*,[17] MD, Cosmo Godino*,[18] MD.

[1] Cardiology Unit, IRCCS Policlinico San Donato, Milan, Italy. Università degli studi di Milano, Milan, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[2] Division of Cardiology, Cardiovascular and Emergency Department, Guglielmo da Saliceto Hospital, Piacenza, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[3] Emergency Department, Ospedali Riuniti San Giovanni di Dio e Ruggi d'Aragona, Salerno, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[4] Cardiology Unit, IRCCS Policlinico San Donato, Milan, Italy. Università degli studi di Milano, Milan, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[5] Università degli studi di Pavia, Pavia, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[6] Università Vita-Salute San Raffaele, Milano, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[7] Division of Cardiology, Cardiovascular and Emergency Department, Guglielmo da Saliceto Hospital, Piacenza, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[8] Division of Cardiology, Cardiovascular and Emergency Department, Guglielmo da Saliceto Hospital, Piacenza, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[9] Division of Cardiology, Cardiovascular and Emergency Department, Guglielmo da Saliceto Hospital, Piacenza, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[10] Division of Cardiology, Cardiovascular and Emergency Department, Guglielmo da Saliceto Hospital, Piacenza, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[11] Division of Cardiology, Cardiovascular and Emergency Department, Guglielmo da Saliceto Hospital, Piacenza, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[12] Clinical Research Unit, Cardiovascular Department, IRCCS Policlinico San Donato, Milan, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[13] Division of Cardiology, Cardiovascular and Emergency Department, Guglielmo da Saliceto Hospital, Piacenza, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[14] Cardiovascular Department, San Raffaele Scientific Institute, Milan, Italy. Università Vita-Salute San Raffaele, Milano, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[15] Cardiology Unit, IRCCS Policlinico San Donato, Milan, Italy. Università degli studi di Milano, Milan, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[16] Arrhythmias and Cardiac Electrophysiology, San Raffaele Scientific Institute, Milan, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[17] Università degli studi di Milano, Milan Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.
[18] Cardiovascular Department, San Raffaele Scientific Institute, Milan, Italy. Università Vita-Salute San Raffaele, Milano, Italy. This author takes responsibility for all aspects of the reliability and freedom from bias of the data presented and their discussed interpretation.

*co-last authorship

Address for correspondence:
Cosmo Godino, MD
Cardio-Thoracic-Vascular Department, San Raffaele Scientific Institute
Via Olgettina 60, 20132 Milan, Italy
Phone: +39022643-3752; Fax:+39022643-7398; email: cosmogodino@gmail.com

Acknowledgement of grant support: none.
The authors report no relationships that could be construed as a conflict of interest.

Key words: acetylsalicylic acid, platelet aggregation inhibitors, COVID-19, SARS-CoV-2, thrombosis.

Background Patients with coronavirus disease 2019 (COVID-19) exhibit high thrombotic risk. The evidence on a potential independent prognostic role of antiplatelet treatment in those patients is limited. The aim of the study was to evaluate the prognostic impact of pre-admission low-dose acetylsalicylic acid (ASA) in a wide series of hospitalized patients with COVID-19.

Methods This cohort study included 984 COVID-19 patients stratified according to ASA intake before hospitalization: ASA^+ (n = 253) and ASA^- (n = 731). Patients were included in ASA^+ group if they received it daily in the 7 days before admission. 213 (83%) were on ASA 100 mg daily. Primary endpoint was a composite of in-hospital death and/or need for respiratory support upgrade, secondary endpoints were in-hospital death and need for respiratory support upgrade.

Results Mean age was 72 [62; 81] with 69% of male patients. ASA^+ patients were significantly older, with higher prevalence of comorbidities. No significant differences regarding the degree of respiratory dysfunction were observed. At 30-day Kaplan-Meier analysis, ASA^+ patients had higher survival free from the primary endpoint and need for respiratory support upgrade, conversely in-hospital death did not significantly differ between groups. At multivariate analysis ASA intake was independently associated with a lower probability of reaching primary endpoint (HR 0.697, 95% C.I. 0.525 – 0.924; $p = 0.012$).

Conclusions In COVID-19 patients undergoing hospitalization, pre-admission treatment with ASA is associated with better in-hospital outcome, mainly driven by less respiratory support upgrade.

Introduction

Coronavirus disease 2019 (COVID-19), caused by the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2),[1] is responsible for the global pandemic outbreak. At the time of this writing, there have been approximately over 180 million cases reported and more than 3.9 million (~2%) deaths due to COVID-19 across more than 200 countries worldwide.[2] Patients with cardiovascular diseases have been reported to have the highest case fatality.[3-4] Although most of COVID-19-related physiopathological pathways remain unclear, some evidences suggest that SARS-CoV-2 infection may predispose patients to thrombosis,[5] both in the arterial and venous circulations,[6] due to inflammation, endothelial dysfunction and, finally, pathological platelet hyperactivation.[7,8] In fact, as Zhang et al. demonstred,[9] SARS-CoV-2 is able to create a spike protein-mediated platelet-ACE2 binding, directly stimulating platelets release of coagulation factors, secretion of inflammatory factors, and formation of leukocyte–platelet aggregates. Furthermore, endothelial cell infection, as evidenced in some autopsy studies,[10,11] or the virus-induced inflammatory response, may contribute to systemic microcirculatory function impairment. The resulting COVID-19-associated endotheliopathy may affect especially, but not only, pulmonary circulation[12] and elicit platelet hyperactivation. For these reasons, antiplatelet therapy, whose impact on outcomes is still under investigation in this subset of patients, may represent an effective therapeutic option.[13-14] Acetylsalicylic acid (ASA) exerts antithrombotic and anti-inflammatory effects, and it had been demonstrated to play some antiviral activity against deoxyribonucleic and ribonucleic acid viruses.[15] The aim of this study was to evaluate the potential protective effect of chronic ASA-based single antiplatelet therapy in a large cohort of patients undergoing hospitalization because of COVID-19.

Methods

This is a multi-center, retrospective, observational study performed at Policlinico San Donato in San Donato Milanese and Ospedale Guglielmo da Saliceto in Piacenza between February 21 and April 22, 2020. The inclusion criteria were: a) patients aged at least 18 years, b) admitted to hospital, c) who were diagnosed COVID-19 according to the interim guidance of the World Health Organization.[16] Clinical information including demographics, comorbidities, medical history, laboratory examinations, baseline and in-hospital treatment measures (including respiratory support) and outcomes was collected after discharge by attending physicians (A.S. and E.P. in San Donato Milanese and A.M. in Piacenza). Each patient underwent admission arterial blood gas analysis, complete blood routine test, including hematologic, biochemical and coagulation function, and chest imaging (X-rays and/or computed tomography) evaluation.

Patients were included in ASA group if they were on treatment and they received it daily at least 7 days before admission.[17,18] ASA treatment was continued during the hospitalization on the same dose as before hospitalization. Patients undergoing orotracheal intubation received ASA by nasogastric tube. Chronic kidney disease (CKD) was defined as Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) equation-derived estimated glomerular filtration rate (eGFR) <60 mL/min/1.73m^2 [19]. The most intense level of oxygen support during hospitalization (nasal cannula, Venturi mask, nonrebreather mask, noninvasive mechanical ventilation [NMV], invasive mechanical ventilation [IMV]) was recorded. According to study institutions' protocols, patients were considered suitable for NMV in the presence of a) moderate-to-high oxygen requirement (partial pressure of oxygen to fractional inspired oxygen ratio [PaO_2/FiO_2] <200 or PaO_2 <60 mmHg or peripheral oxygen saturation <94% or 88% in patients with acute or acute on chronic type II respiratory failure, despite 15 L/min oxygen administration via nonrebreather mask), b) in the absence of contraindication to using NMV. IMV was considered after unsuccessful NMV, defined as PaO_2/FiO_2 tending to decrease and PaO_2 <60 mmHg or if NMV was not advisable, if patient clinical status allowed. Patients requiring IMV at the time of admission were not included in

the study because a) only in-hospital death, but not primary endpoint, could have been evaluated since IMV represent the most intense level of respiratory support, and b) those patients belong to critically-ill category, in which the underlying thrombotic and inflammatory damage may have been too advanced to have been influenced by ASA intake. Finally, because of different mechanism of action, P2Y12 inhibitors-assuming patients (i.e. clopidogrel) were not considered in our analysis.

The primary study endpoint was a composite of 30-day in-hospital death or need for respiratory support upgrade to NMV, including Continuous Positive Airway Pressure (CPAP) and Bilevel positive airway pressure (BiPAP) or IMV. The secondary clinical endpoints were in-hospital death and need for respiratory support upgrade up to 30-day, analyzed individually.

This study complied with the principles outlined in the 1975 Declaration of Helsinki. The study was approved by the Local Ethics Committee.

Given the retrospective nature of our study, no statistical sample size calculation was performed a priori, and sample size was equal to the number of eligible patients hospitalized during the study period. Distribution of continuous data was tested with the Shapiro-Wilk test. Non-normally distributed variables were presented as median and interquartile range. Categorical variables were reported as absolute numbers and percentages. Continuous variables were then compared using Mann-Whitney U test; categorical variables were compared with Chi square test. Event-free survival up to 30-day were evaluated according to the unadjusted Kaplan-Meier method and survivals among groups were compared using log-rank test (Cox-Mantel test). Cox proportional hazards regression analysis was used to determine significant predictors of primary and secondary endpoints. Variables with a univariate statistical significance of <0.05 were selected for inclusion into the multivariable model. Multivariate analysis, using stepwise forward selection, was finally performed to analyze the association of baseline characteristics with study endpoints, expressed as hazard ratio (HR) with 95% confidence interval (CI) and *p* values. All statistical tests were 2-sided, and *p* values <0.05 were considered statistically significant. The statistical analyses were performed

using SPSS software version 25.0.0 (SPSS Inc., Chicago, IL) and GraphPad Prism software version 6 (GraphPad, Inc, San Diego, CA).

Results

Nine hundred and eighty-four patients (200 in San Donato Milanese and 784 in Piacenza) with COVID-19 (median age 72 [62; 81] years; 69% male) were included in the study, **Figure S1, Supplementary material**. According to baseline pre-admission ASA intake we identified two groups, 253 (26%) patients were receiving ASA (ASA^+) and 731 were not (74%) (ASA^-). Concerning ASA^+ patients, 213 (83%) were on ASA 100 mg daily, meanwhile the remaining were assuming it a daily dose of 75 mg.

Compared to ASA^- patients, the group ASA^+ was significantly older and suffered more from cardiovascular comorbidities, such as hypertension, diabetes mellitus and dyslipidemia, resulting in higher incidence of coronary artery disease, peripheral artery disease and previous ischemic stroke or transient ischemic attack. Heart failure, chronic kidney disease and chronic obstructive pulmonary disease were less frequent in ASA^+ group. ASA^+ patients were more often on angiotensin-converting enzyme inhibitors or angiotensin type 1 receptor blocker and statin therapy. Of note, there were no differences either on pre-hospitalization infection-related symptoms, except for lower incidence of fever in ASA^+ group, or on time between symptoms onset and hospitalization. Arterial blood gas analysis at admission showed similar degree of respiratory impairment. Notably, 32 patients required NMV at the time of admission, without significant differences between groups (p = 0.158). Besides, ASA^- patients presented with significantly higher neutrophils to lymphocytes ratio (N/L, 5 [3; 9] vs. 4 [2; 7], p = 0.013) and hemoglobin levels (14 [12; 15] g/dL vs. 13 [12; 15] g/dL, p = 0.016). ASA^+ patients showed a worse baseline renal function, as assessed by lower median estimated glomerular filtration rate (eGFR, 59 [43; 80] mL/min/1.73 m^2 vs. 70 [51; 89] mL/min/1.73 m^2, p = 0.001) and increased high-sensitivity troponin T values (24 [14; 65] ng/L vs. 12 [7; 24] ng/L, p <0.001), whereas liver function indexes did not differ between study groups. Serum D-dimer level was similar among two groups (2.01 [0.90; 3.53] µg/mL vs. 1.50 [0.69; 3.0] µg/mL, p = 0.276). Chest imaging revealed bilateral interstitial infiltrates in 90% of entire study cohort, without significant difference between study groups (p = 0.774).

Admission risk scores assessing in-hospital mortality did not differ significantly between groups. During hospitalization empirical anti-SARS-CoV-2 therapy, including tocilizumab, antibiotic, glucocorticoid and low-molecular weight heparin (LMWH), was administered more often to ASA^- patients, as well as oxygen therapy, given that ASA^+ patients underwent less NMV or IMV treatments (p <0.001), **Table 1**.

Median length of in-hospital stay was 11 [7; 18] days, similar between the two groups (p = 0.980). Furthermore, no significant differences were observed in patients assuming ASA 100 mg daily compared to those taking 75 mg daily (p = 0.331) concerning duration of hospitalization. At 30-day Kaplan Meier analysis in the entire study cohort, compared to ASA^- patients, ASA^+ suffered less adverse events in terms of both primary endpoint (63% vs. 75%; HR 0.788, log-rank p = 0.013) and need for respiratory support upgrade (33% vs. 49%; HR 0.640, log-rank p = 0.008), **Figure 1, panel A and C**, respectively, with 19% ASA^+ patients vs. 25% ASA^- patients needing upgrade to NMV (log-rank p = 0.006), and 15% ASA^+ patients vs. 25% ASA^- patients needing upgrade to IMV (log-rank p = 0.017). Meanwhile in-hospital death did not differ significantly between two groups (ASA^+ 52% vs. ASA^- 53%; HR 1.042, log-rank p = 0.653), **Figure 1, panel B.** Primary and secondary endpoints are shown in **Table S1, Supplementary material**.

At univariate Cox regression analysis, ASA, as well as glucocorticoid therapy, was associated with better outcome in terms of primary endpoint, whereas age, male gender, hypertension, admission N/L >3 and eGFR <60 mL/min/1.73m^2 correlated with a worse one. Multivariate analysis identified ASA use as an independent positive prognostic factor in terms of primary endpoint (HR 0.697, 95% C.I. 0.525 – 0.924; p = 0.012), **Figure 2**. ASA was also identified as independent protective factor in terms of need for less respiratory support upgrade (HR 0.529, 95% C.I. 0.333 – 0.839; p = 0.007), meanwhile it was not able to predict in-hospital death, as evidenced in **Table 2.**

Finally, Cox regression analysis showed no significant impact of different doses of ASA in terms of primary endpoint (HR 0.769, 95% C.I. 0.489 – 1.209, p = 0.256), in-hospital death (HR 0.864, 95%

C.I. 0.525 – 1.422, $p = 0.565$) and need for respiratory support upgrade (HR 0.828, 95% C.I. 0.368 – 1.865, $p = 0.649$).

Discussion

The cardinal finding of this multi-center, observational analysis with the prespecified hypothesis of a protective role of ASA in COVID-19 infection was that pre-admission chronic ASA therapy resulted in a better in-hospital outcome mainly driven by less respiratory support upgrade. This noteworthy finding was associated with no difference concerning in-hospital death among patients with or without ASA.

An intriguing question involving the scientific community is the definition of the role played by antithrombotic treatment in COVID-19 patients,[7,20] primarily focusing on anticoagulation and its clinical impact.[21-22] Considering the lack of a standard of care, dominant related questions are: 1) what is the best antithrombotic strategy (anticoagulant with or without antiplatelet and eventually which specific drug)? and, 2) which kind of clinical benefit to expect from, and primarily which kind of benefit to consider as still useful (freedom from complications and/or survival improvement) for each patient within the broad spectrum of presentation?

The present study is an attempt to provide some potential answers and to make a firm focus on the role of ASA, a therapeutic regimen approved in patients phenotype with multiple cardiovascular comorbidities and more exposed to COVID-19 injury. To the best of our knowledge, this is the largest analysis showing an association between ASA and favourable outcome in COVID-19 patients. The present findings are consistent with a multi-centre study, where ASA was independently associated with decreased risk of mechanical ventilation, intensive care unit admission and finally in-hospital mortality, though in a smaller sample size (412 patients).[23] Conversely, in our analysis ASA failed to predict overall survival. Apparently divergent results may be associated to either patient selection resulting in different baseline clinical features or different level of adjustment for several prognostic confounders. However, since a sub-analysis of the TARGET-COVID study showed an insufficient pharmacodynamic effect of 81 mg daily ASA in a high percentage of COVID-19 patients, most of whom African Americans,[24] it is at least surprising how low-dose (median 81 mg daily) ASA is sufficient to provide such a meaningful clinical effect.

Our results suggest that, although suffering from a similar extent of disease, ASA- patients underwent in-hospital progressive clinical deterioration and were in greater need of empirical anti-SARS-CoV-2 therapy and respiratory support, potentially related to a pathological platelet hyperactivation. Through inhibition of synthesis of cyclooxygenase and activation of nuclear factor-κB,[13] ASA exerts a simultaneous antiplatelet and anti-inflammatory effect, potentially able to prevent intravascular coagulation and neutrophil-mediated microvascular thrombosis, as showed in animal model.[25] Since platelets may represent a bridge between immune system and thrombosis, therefore the frontline of COVID-19 pathogenesis,[26] antiplatelet therapy may constitute a cost-effective, relatively low risk-associated,[27] therapeutic strategy to prevent patients from clinical worsening during SARS-CoV-2 infection in addition to LMWH, especially in non-critically ill patients. Indeed, our analysis identified LMWH as an independent predictor of in-hospital mortality. That is consistent with recently published data deriving from a single multiplatform, randomized, controlled trial suggesting that in the moderately ill patients therapeutic-dose LMWH appeared to increase the probability of survival until hospital discharge.[28] Furthermore, the preprint article reporting the findings of the Therapeutic Anticoagulation versus Standard Care as a Rapid Response to the COVID-19 Pandemic (RAPID) trial showed as therapeutic anticoagulation group had a lower incidence of death at 28 days.[29]

Therefore, it is reasonable that, rather than a single medication, combination therapies targeting several pathological pathways (e.g., inflammation, coagulopathy, thrombocytopathy and endotheliopathy) are more likely to be successful.

The identification of single-patient thrombogenic phenotype, based upon not only thrombotic biomarkers such as D-dimer, fibrinogen, prothrombin time and activated partial thromboplastin time, but also whole blood viscoelastic analysis performed by thromboelastography or rotational thromboelastometry, would allow personalizing antithrombotic therapy in COVID-19 patients and possibly improve outcomes.[30] Interestingly, as Gurbel et al. suggested,[31] systemic concentrations of oral-administered ASA may not reach the airway and alveolus to effectively reduce the virus

load. In this perspective, unconventional routes of administration, including inhaled nanoparticle, should be considered to achieve locally effective concentrations.

To date available data are not sufficient to influence standard of care. Randomized controlled trial, such as the ongoing Randomized Evaluation of COVID-19 Therapy (RECOVERY) trial (NCT04381936), will definitively evaluate whether antiplatelet therapy prevents adverse outcome in patients with COVID-19.

The present study suffers from the following limitations. In view of the observational nature of our analysis, patient selection and ascertainment bias may have influenced event rates. Particularly, identifying study groups according to pre-admission ASA intake represents a selection bias, since patients were on treatment because of the presence of more cardiovascular comorbidities. Furthermore, we did not account for safety endpoints, such as major bleeding.

Accordingly, our results should be considered as hypothesis generating and need confirmation in further larger observational studies or randomized trials.

In conclusion, in this retrospective analysis of patients with COVID-19 undergoing hospitalization, ASA is associated with a better in-hospital outcome in terms of in-hospital death or need for respiratory support upgrade, whose definitive evidence is mainly supported by the latter.

Acknowledgments: all authors have participated in the work and have reviewed and agree with the content of the article.

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Figure titles and legends

Figure 1. Entire study cohort 30-day Kaplan-Meier analysis of primary and secondary endpoints.

Entire study cohort 30-day Kaplan-Meier analysis of survival free from *primary endpoint* (panel A), *in-hospital death* (panel B) and *need for respiratory support upgrade* (panel C).

ASA: acetylsalicylic acid.

Figure 2. Forest plot showing results from multivariate Cox regression analysis regarding primary endpoint.

ASA: acetylsalicylic acid. eGFR: estimated glomerular filtration rate. N/L: neutrophils to lymphocytes ratio.

Data are presented as hazard ratio (HR) with 95% confidence interval (CI).

Table 1. Baseline clinical features, in-hospital instrumental evaluation and empirical therapy in entire study cohort and the two subgroups identified according to baseline acetylsalicylic acid intake.

	entire study cohort (n = 984)	**ASA$^+$ (n = 253)**	**ASA$^-$ (n = 731)**	***p* value**
Clinical features on admission				
Age (years)	72 [62; 81]	76 [67; 82]	71 [61; 80]	**<0.001**
Male gender, n (%)	678 (69)	171 (68)	507 (69)	0.601
Initial common symptoms[a]				
Fever, n (%)[c]	644 (91)	165 (86)	479 (93)	**0.006**
Dry cough, n (%)	369 (56)	91 (50)	278 (58)	0.112
Dyspnea, n (%)	449 (67)	129 (69)	320 (66)	0.401
Diarrhea, n (%)	60 (10)	11 (6)	49 (11)	0.094
Symptoms onset to admission (days)	7 [4; 10]	7 [3; 9]	7 [4; 10]	0.242
Comorbidities, n (%)[b]				
Hypertension	604 (62)	215 (85)	389 (54)	**<0.001**
Diabetes mellitus	188 (19)	85 (34)	103 (14)	**<0.001**
Dyslipidemia	237 (24)	112 (44)	125 (17)	**<0.001**
Coronary artery disease	86 (9)	81 (33)	21 (3)	**<0.001**
Heart failure	95 (10)	58 (23)	37 (5)	**<0.001**
Atrial fibrillation	124 (13)	31 (12)	93 (13)	0.774
Peripheral artery disease	30 (3)	15 (6)	15 (2)	**0.002**
Previous ischemic stroke/TIA	31 (3)	13 (5)	18 (3)	**0.041**
Chronic kidney disease	98 (10.1)	42 (17)	56 (8)	**<0.001**
Chronic obstructive pulmonary disease	140 (14)	53 (21)	87 (12)	**0.001**
History of neoplasia	60 (6)	19 (8)	41 (6)	0.307
Drugs, n (%)[b]				
Anticoagulant				0.402
OAT	68 (7)	19 (8)	49 (7)	
DOAC	45 (5)	7 (3)	38 (5)	
ACE-I/ARB[d]				**<0.001**

ACE-I	250 (26)	90 (36)	160 (23)	
ARB	180 (19)	73 (29)	107 (15)	
Statin	208 (27)	100 (48)	108 (19)	**<0.001**
Vital signs				
Systolic blood pressure (mmHg)	130 [117; 145]	130 [115; 145]	130 [120; 145]	0.888
Diastolic blood pressure (mmHg)	75 [70; 80]	70 [65; 80]	77 [70; 83]	**0.017**
Heart rate (bpm)	90 [80; 100]	87 [76; 100]	90 [80; 100]	0.097
Respiratory rate (min^{-1})	22 [18; 25]	22 [18; 25]	22 [18; 25]	0.498
Body temperature (°C)	38 [37; 38]	37.7 [37; 38]	38 [37; 38.5]	**0.027**
Peripheral oxygen saturation (%)	91 [87; 94]	91 [87; 94]	91 [87.5; 94]	0.995
Arterial blood gas				
pH	7.47 [7.43; 7.50]	7.47 [7.43; 7.51]	7.47 [7.43; 7.50]	0.964
PaO_2 (mmHg)	60 [50; 70]	59 [50; 67]	60 [50; 71]	0.815
PaO_2/FiO_2 (mmHg/%)	2.81 [2.33; 3.20]	2.79 [2.29; 3.13]	2.81 [2.34; 3.28]	0.662
$PaCO_2$ (mmHg)	33 [30; 37]	33 [29; 37]	30 [30; 36]	0.212
HCO_3^- (mmol/L)	24 [22; 27]	25 [21; 28]	24 [22; 26]	0.518
SO_2 (%)	94 [91; 96]	93 [91; 94]	94 [91; 96]	0.320
Lactate (mmol/L)	1.3 [0.9; 1.8]	1.3 [0.7; 1.8]	1.3 [0.9; 1.7]	0.689
Laboratory indices				
White blood cells (10^9/L)	6.9 [5.2; 9.5]	6.5 [4.8; 8.6]	7.1 [5.3; 9.8]	0.061
Neutrophils (10^9/L)	5.0 [3.1; 7.9]	4.6 [2.8; 7.1]	5.2 [3.2; 8.1]	0.101
Lymphocytes (10^9/L)	1.2 [0.8; 1.6]	1.2 [0.8; 1.8]	1.1 [0.8; 1.6]	**0.043**
N/L	5 [2; 9]	4 [2; 7]	5 [3; 9]	**0.013**
Hemoglobin (g/dL)	14 [12; 15]	13 [12; 15]	14 [12; 15]	**0.016**
Hematocrit (%)	41 [37; 44]	40 [36; 44]	41 [37; 45]	0.104
Platelets, (10^9/L)	200 [150; 263]	206 [140; 282]	195 [153; 260]	0.125
Creatinine (mg/dL)	1.0 [0.9; 1.3]	1.1 [0.9; 1.43]	1.0 [0.8; 1.3]	0.135
eGFR (mL/min/1.73 m^2)	68 [48; 87]	59 [43; 80]	70 [51; 89]	**0.001**
Urea (mg/dL)	45 [32; 64]	48 [35; 68]	44 [31; 63]	0.097
Sodium (mEq/L)	137 [134; 139]	137 [134; 140]	137 [134; 139]	0.806

Potassium (mEq/L)	4.14 [3.80; 4.53]	4.27 [3.80; 4.70]	4.10 [3.80; 4.50]	0.172
Lactate dehydrogenase (UI/L)	451 [344; 588]	443 [323; 580]	455 [351; 592]	0.484
Creatinine kinase (UI/L)	119 [64; 259]	126 [68; 257]	118 [63; 26]	0.579
Total bilirubin (mg/dL)	0.69 [0.51; 0.93]	0.70 [0.51; 0.96]	0.69 [0.52; 0.91]	0.654
Glutamic pyruvic transaminase (UI/L)	30 [20; 49]	29 [18; 47]	30 [21; 50]	0.857
Glutamic oxaloacetic transaminase (UI/L)	42 [30; 66]	42 [30; 67]	42 [31; 65]	0.298
High-sensitivity troponin T (ng/L)[e]	16 [8; 30]	24 [14; 65]	12 [7; 24]	**<0.001**
C-reactive protein (mg/dL)	10 [5; 16]	9 [5; 15]	10 [5; 17]	0.103
Serum ferritin (ng/mL)	876 [510; 1460]	715 [446; 1601]	940 [529; 1460]	0.120
D-dimer (µg/mL)	1.53 [0.77; 3.18]	2.01 [0.90; 3.53]	1.50 [0.69; 3.0]	0.276
Fibrinogen (mg/dL)	602 [486; 734]	632 [485; 718]	595 [482; 739]	0.811
Chest imaging, n (%)				
Bilateral infiltrates	886 (90)	228 (90)	658 (90)	0.774
Pleuric effusion	177 (18)	66 (26)	110 (15)	**0.004**
Risk scores				
qSOFA	1 [0; 1]	1 [0; 1]	1 [0; 1]	0.226
CURB-65	1 [0; 2]	1 [0; 1.75]	1 [0; 1]	0.394
Drugs, n (%)[a]				
Hydroxychloroquine	494 (78)	127 (76)	367 (74)	0.323
Tocilizumab	57 (16)	4 (4)	53 (20)	**<0.001**
Antibiotic[f]	439 (69)	102 (60)	337 (73)	**0.001**
Glucocorticoid	207 (34)	43 (27)	164 (37)	**0.019**
Low-molecular weight heparin				**0.015**
none	246 (40)	79 (47)	167 (37)	
prophylactic dose	39 (6)	14 (9)	25 (6)	
therapeutic dose	331 (54)	74 (44)	257 (57)	
Oxygen therapy, n (%)[a]				**<0.001**
None	193 (20)	35 (14)	158 (22)	
Nasal cannula/Venturi mask/nonrebreather mask	454 (48)	146 (59)	308 (44)	
Noninvasive ventilation	184 (19)	41 (17)	143 (20)	

Invasive mechanical ventilation	123 (13)	25 (10)	98 (14)

ACE-I: angiotensin-converting enzyme inhibitors. ARB: angiotensin 1 receptor blocker. ASA: acetylsalicylic acid. DOAC: direct oral anticoagulant. eGFR: estimated glomerular filtration rate. HCO_3^-: hydrogen carbonate. N/L: neutrophils to lymphocytes ratio. NT-proBNP: N-terminal prohormone of brain natriuretic peptide. OAT: oral anticoagulant therapy. PaO_2: partial pressure of oxygen. PaO_2/FiO_2: partial pressure of oxygen to fractional inspired oxygen ratio. qSOFA: quick sepsis related organ failure assessment. SO_2: oxygen saturation. TIA: transient ischemic attack.

Data are presented as n (%), mean ± SD or median [IQR].

[a]Values are avaible for ~ 70% of the entire study cohort.

[b]Values are avaible for ~ 97% of the entire study cohort.

[c]Fever was classified as highest patient temperature 37.3 °C or higher. To minimize interference of treatment, the highest patient temperature was defined using the self-reported highest temperature before taking antipyretic drug.

[d]ACE-I/ARB use was defined as use of these drugs at the time of admission that continued through hospitalization.

[e]Values are avaible for ~ 25% of the entire study cohort.

[f]Including azithromycin 500 mg daily dose p.o. and/or ceftriaxone 2000 mg daily dose i.v.

Table 2. Primary endopoint-related univariate and multivariate Cox regression analysis in entire study cohort.

	Univariate analysis			Multivariate analysis		
	HR	95% CI	*P* value	HR	95% CI	*P* valu
hospital death and/or piratory support upgrade						
	1.024	1.016 – 1.031	<0.001			
e gender	1.312	1.085 – 1.585	0.005	1.424	1.100 – 1.842	0.007
ertension	1.244	1.039 – 1.490	0.018			
A	0.788	0.647 – 0.960	0.018	0.697	0.525 – 0.924	0.012
>3	1.549	1.239 – 1.938	<0.001	1.483	1.145 – 1.919	0.003
FR <60 mL/min/1.73m^2	1.466	1.207 – 1.780	<0.001	1.351	1.054 – 1.731	0.018
cocorticoid	0.698	0.558 – 0.872	0.002	0.782	0.622 – 0.985	0.036
hospital death						
	1.069	1.058 – 1.081	<0.001	1.066	1.047 – 1.085	<0.001
ertension	1.577	1.251 – 1.987	<0.00			
rt failure	1.728	1.275 – 2.342	<0.001			
vious ischemic stroke/TIA	1.619	1.008 – 2.601	0.046			
onic obstructive pulmonary ease	1.411	1.069 – 1.862	0.015			
>3	1.519	1.128 – 2.045	0.006	1.468	1.054 – 2.045	0.023
moglobin	0.913	0.855 – 0.975	0.007			
FR <60 mL/min/1.73m^2	2.693	2.071 – 3.502	<0.001	1.728	1.263 – 2.365	0.001
-molecular weight heparin	0.640	0.477 – 0.858	0.003	0.660	0.487 – 0.893	0.007
piratory support upgrade						
e gender	2.084	1.473 – 2.949	<0.001	1.855	1.232 – 2.794	0.003
A	0.640	0.458 – 0.894	0.009	0.529	0.333 – 0.839	0.007
>3	1.706	1.211 – 2.404	0.002			
-molecular weight heparin	1.652	1.154 – 2.364	0.006			
cocorticoid	0.477	0.346 – 0.659	<0.001	0.556	0.395 – 0.782	0.001

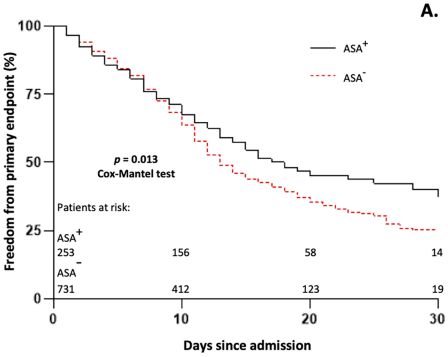
ASA: acetylsalicylic acid. eGFR: estimated glomerular filtration rate. N/L: neutrophils to lymphocytes ratio. TIA: transient ischemic attack.

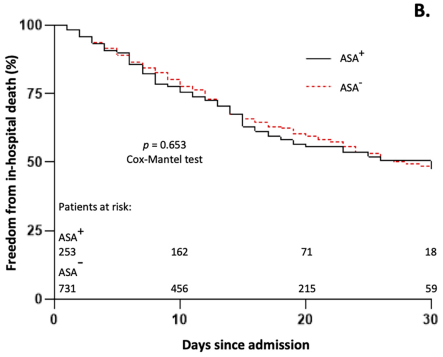
Data are presented as hazard ratio (HR) with 95% confidence interval (CI) and *p* values. Only covariates with a univariate statistical significance of <0.05 were reported.

CRediT author statement

Antonio Sisinni: Formal analysis, Writing - Original Draft, Visualization
Luca Rossi: Investigation
Antonio Battista: Conceptualization, Writing - Review & Editing
Enrico Poletti: Investigation
Federica Battista: Formal analysis
Rosa Alessia Battista: Formal analysis
Alessandro Malagoli: Investigation
Andrea Biagi: Investigation
Alessia Zanni: Investigation
Concetta Sticozzi: Investigation
Greta Comastri: Investigation
Massimiliano M Marrocco-Trischitta: Supervision
Alberto Monello: Investigation
Alberto Margonato: Supervision
Francesco Bandera: Writing - Review & Editing
Pasquale Vergara: Supervision
Marco Guazzi: Writing - Review & Editing
Cosmo Godino: Conceptualization, Methodology, Writing - Review & Editing, Visualization

- Coronavirus disease 2019 (COVID-19) may predispose patients to thrombosis.
- Antiplatelet therapy may represent an effective therapeutic option.
- Patients on pre-admission acetylsalicylic acid therapy suffered less from in-hospital death and/or need for respiratory support upgrade.


A.
Freedom from primary endpoint (%)
100
75
50
25
0
ASA+
ASA-
p = 0.013
Cox-Mantel test
Patients at risk:
ASA+
253 156 58 14
ASA-
731 412 123 19
0 10 20 30
Days since admission


B.
Freedom from in-hospital death (%)
100
75
50
25
0
ASA+
ASA-
p = 0.653
Cox-Mantel test
Patients at risk:
ASA+
253 162 71 18
ASA-
731 456 215 59
0 10 20 30
Days since admission

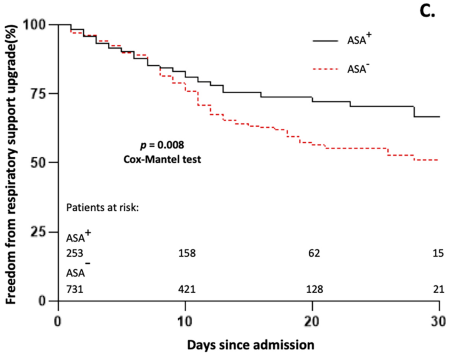
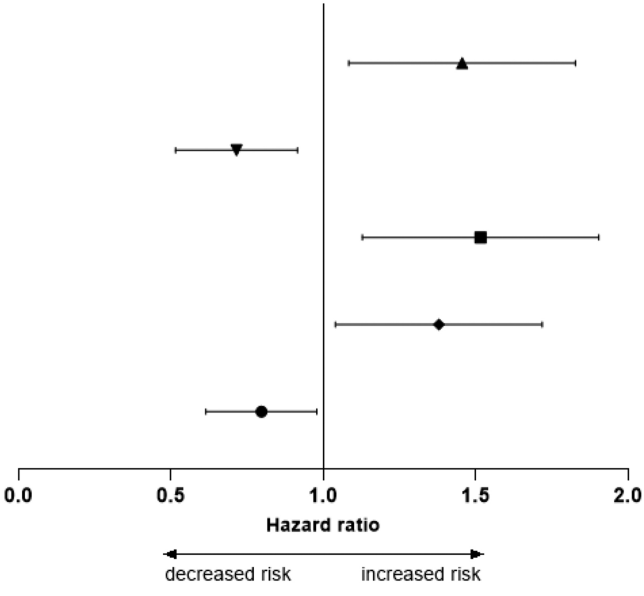

C.
Freedom from respiratory support upgrade(%)
100
75
50
25
0
ASA+
ASA-
p = 0.008
Cox-Mantel test
Patients at risk:
ASA+
253 158 62 15
ASA-
731 421 128 21
0 10 20 30
Days since admission

Figure 1



	HR [95% CI]
male gender	1.424 [1.100 – 1.842]
ASA	0.697 [0.525 – 0.924]
N/L >3	1.483 [1.145 – 1.919]
eGFR <60 mL/min/m^2	1.351 [1.054 – 1.731]
glucocorticoid	0.782 [0.622 – 0.985]

Figure 2